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<b>(21) International Application Number:</b> PCT/US92/07029 <b>(22) International Filing Date:</b> 20 August 1992 (20.08.92)  <b>(30) Priority data:</b> 747,371 20 August 1991 (20.08.91) US  <b>(71) Applicant:</b> THE UNITED STATES OF AMERICA, represented by THE SECRETARY, DEPARTMENT OF HEALTH AND HUMAN SERVICES [US/US]; National Institutes of Health, Office of Technology, Transfer, Box OTT, Bethesda, MD 20892-9902 (US).  <b>(72) Inventor:</b> CRYSTAL, Ronald, G. ; 13712 Canal Vista Court, Potomac, MD 20854 (US).		<b>(74) Agents:</b> GAGALA, Bruce, M. et al.; Leydig, Voit & May- er, Two Prudential Plaza, Suite 4900, Chicago, IL 60601-6780 (US).  <b>(81) Designated States:</b> AU, CA, JP, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, SE).  <b>Published</b> <i>With international search report.</i>
<b>(54) Title:</b> ADENOVIRUS MEDIATED TRANSFER OF GENES TO THE GASTROINTESTINAL TRACT  <b>(57) Abstract</b>  The present invention relates, in general, to an adenovirus mediated transfer of genes to the gastrointestinal tract. In parti- cular, the present invention relates to a method of recombinant, replication-deficient adenovirus mediated transfer of therapeutic genes to the gastrointestinal tract whereby therapeutic proteins for systemic and/or local purposes are produced.		

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**ADENOVIRUS MEDIATED TRANSFER  
OF GENES TO THE GASTROINTESTINAL TRACT**

Field of the Invention

The present invention relates, in general,  
5 to a method of adenovirus mediated transfer of  
genes to the gastrointestinal tract. In  
particular, the present invention relates to a  
method of recombinant, replication-deficient  
adenovirus mediated transfer of therapeutic genes  
10 to the gastrointestinal tract for the purpose of  
producing therapeutic proteins for systemic  
and/or local use.

Background Information

The use of proteins as therapeutic agents  
15 is limited inter alia by the physiologic barrier  
of the gastrointestinal tract. The terms  
protein, polypeptide, peptide, or segment of  
amino acids are herein used interchangeably to  
define a polymer of amino acids linked through  
20 peptide bonds. Therapeutic proteins are defined  
herein as proteins advantageous to an individual.  
Proteins cannot be administered for therapeutic  
purposes by the oral or rectal routes because  
they will not generally reach the circulation in  
25 an intact form in concentrations needed for  
therapy (the proteins are degraded and/or not  
absorbed). Consequently, therapeutic proteins  
need to be administered systemically for example,  
by the intravenous, subcutaneous, intradermal or  
30 intramuscular routes.

This problem of the administration has  
been dramatically heightened by the development  
of recombinant DNA technology, where it is  
possible to produce many different therapeutic

proteins, all of which have to be administered systemically. While this may not be a major problem for short term use, long term use (which is the typical use for most of the recombinant proteins) requires long term systemic administration with all of the attendant problems with access route (e.g., veins available, discomfort and cost).

It is known that recombinant adenoviruses can be used to produce human protein in vivo (examples include injection of recombinant adenovirus intravenously into the portal vein to the liver, and intratracheal to the lung. All publications mentioned herein are hereby incorporated in their entirety by reference. (see Rosenfeld M et al. (1991) Science 252:431-434; Jaffe HA et al. (1991) Clin Res 39(2) 302A; Rosenfeld MA et al. (1991) Clin Res 39(2): 311A). However, all of these approaches are impractical to use for systemic administration of recombinant proteins because they require parenteral administration of the recombinant gene (i.e., intravenous, intraportal, intratracheal).

The present invention circumvents this by providing a method of administering therapeutic proteins by enteral routes by using a recombinant, replication deficient adenovirus containing the coding sequences of the gene of the therapeutic protein to insert the gene into the lining cells of the gastrointestinal tract, and using that site to produce the protein and secrete it into the circulation where the therapeutic protein would be available for systemic use. As an alternative, the same approach can be used to secrete proteins into the gastrointestinal tract for local therapeutic use within the lumen of the gastrointestinal tract, or for

use within the cells or extracellular matrix of the walls of the gastrointestinal tract.

Studies in the 1960's demonstrated that live adenovirus placed into enteric coated capsules (to avoid inactivation in the stomach) and administered to humans by the oral route resulted in systemic immunization against the adenovirus (Chanock RM et al. (1966) JAMA 195:151-158). This is now a standard immunization procedure against adenovirus for military recruits in the USA. The concept underlying this immunization strategy is that the adenovirus will leave the capsule as it dissolves in the lumen of the intestine, infect the intestinal epithelial cells, replicate in the epithelial cells and the resulting shed newly replicated virus presents itself to the immune system, resulting in systemic immunity against the adenovirus.

Previously, it has been demonstrated that the adenovirus can be modified so that it is replication deficient (i.e., will not direct the production of new virus after it infects its target cell) and so that it contains new genes (e.g., the coding sequences of human genes of therapeutic interest). The use of recombinant DNA inserts under the direct control of the early promoter (EP) of the Ela region of the adenovirus genome has been described by M. Perricaudet, et al. of the Pasteur Institute in European Patent Application No. 0185573, published June 25, 1986. Such a modified virus can be used to transfer the recombinant gene to target cells in vivo (for examples, see Rosenfeld M et al. (1991) Science 252:431-434; Berkner KL (1988) BioTechniques 6:616-629).

### SUMMARY OF THE INVENTION

It is a general object of this invention to provide a method of producing a protein in the cells of the gastrointestinal tract of a patient.

5 It is a specific object of this invention to provide a method of producing a protein in the cells of the gastrointestinal tract of a patient. The method comprises administering to the patient's gastrointestinal tract a replication  
10 deficient adenovirus comprising a DNA segment encoding the protein under conditions such that the protein is produced. Depending on the specific sequences placed into the recombinant adenovirus, the protein would preferably then be  
15 secreted for systemic therapy to the circulation, for local therapy to the lumen of the gastrointestinal tract, or both. Further, the design of the recombinant adenovirus may preferably be to deliver the protein for use  
20 within the cells of the gastrointestinal tract or in the walls of the gastrointestinal tract.

Further objects and advantages of the present invention will be clear from the description that follows.

### 25 BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1. Recombinant adenovirus (Ad) vector. Top - wild type Ad5 genome showing the E1a, E1b [map units (mu) 1.3-11.2; 100 mu = 36 kb] and E3 (mu 76.6-86.0) regions.

30 Figure 2. Anatomy of the colon wall and the cultured epithelial cell model used to evaluate the polarity of secretion of  $\alpha$ 1-antitrypsin produced by T84 human colon carcinoma epithelial cells modified with the recombinant  
35 adenovirus Ad- $\alpha$ 1AT (ATCC CCL 248).

A. Cross-section of the colon wall showing the epithelial cells (14), the lumen of the colon (13), the apical surface (12) of the epithelium abutting the lumen, the basolateral surface (11) of the epithelium abutting the submucosa and thus the capillaries (19) and the muscle layer (20).

B. Chamber for epithelial cell cultures showing the microporous membrane (17), the cultured cells (16), and the separated apical (15) and basolateral (18) compartments.

Figure 3. Demonstration of de novo synthesis and secretion of human  $\alpha$ 1-antitrypsin ( $\alpha$ 1AT) by rat colon exposed to the recombinant adenovirus Ad- $\alpha$ 1AT ex vivo.

#### DETAILED DESCRIPTION OF THE INVENTION

The present invention relates to a method of producing therapeutic proteins in the gastrointestinal tract. As a step, first, a replication deficient adenovirus (referred to below as the "modified adenovirus") is constructed with the coding sequences of the protein of therapeutic interest. As an example, the adenovirus Ad- $\alpha$ 1AT containing the coding sequence of the human  $\alpha$ 1AT gene is used (see Figure 1 and Rosenfeld M et al. (1991) Science 252:431-434). Second, the modified adenovirus is placed into an enteric capsule (or alternatively, administered via tube past the stomach, orally or by tube into the stomach after the stomach lining fluid has been modified such that the virus will not be altered; or via the rectal route). Alternatively, a special coating may be applied to the adenovirus to prevent release and absorption of the modified adenovirus until the tablet reaches the basic (pH) environment of the duodenum, jejunum, ileum, or colon. For oral

administration a capsule tablet or pill are convenient delivery vehicles; for rectal administration, a suppository may be preferred. Administration is then effected. The following scenario then occurs: (1) the cells (preferably, the epithelial cells) of the gastrointestinal tract are infected by the modified adenovirus; (2) the recombinant gene in the modified adenovirus sequence directs the synthesis of the recombinant protein which (depending upon how the sequences in the recombinant gene are engineered) can then be secreted into the circulation, into the lumen of the gastrointestinal tract or into both the circulation and the lumen of the gastrointestinal tract or within the epithelial cells of the gastrointestinal tract or in the local environs within the wall of the gastrointestinal tract; and (3) the therapeutic protein is then available to act systemically (when secreted into the circulation) or in the intestine (when secreted into the lumen), within the cells and/or the extracellular matrix of the wall of the gastrointestinal tract.

The present invention provides a practical, easy and safe way to administer recombinant proteins to humans. By using a replication deficient adenovirus, the process is safe because the virus cannot replicate in the target cells. By using a recombinant adenovirus, the target cells will produce the human therapeutic protein. By choosing the epithelium of the gastrointestinal tract for the target of infection by the replication deficient recombinant adenovirus, the invention permits ease of administration (preferably, by oral route via enteric coated capsule) via a route that can be used repetitively (for example, daily, or less



frequently, depending on the chronicity of the recombinant adenovirus infection in the epithelial cells) and safely.

Because the epithelial cells of the gastrointestinal tract will secrete some of the product of the recombinant adenovirus through the basolateral surface of the infected epithelial cells, the method is available for applications requiring systemic use. Because these epithelial cells also secrete some of the product through their apical surface, the method is available for applications requiring luminal use (for example, intraluminal gastrointestinal disorders and gastrointestinal cancer). If the recombinant adenovirus is designed appropriately, the therapeutic protein will be available for therapeutic use within the epithelial cells of the gastrointestinal tract or in the local environs of the wall of the gastrointestinal tract (for example, for gastrointestinal tract cancer or gastrointestinal inflammatory disorders).

For disorders requiring systemic administration, this approach provides an easy and safe manner of administering recombinant protein to the circulation. Examples of such proteins include, but are not limited to:

- $\alpha$ 1-antitrypsin - for  $\alpha$ 1-antitrypsin deficiency
- factor VIII - for hemophilia
- other coagulation factors - for bleeding disorders
- growth hormone - for growth disorders

- insulin - for diabetes
- other peptide hormones
- other pituitary hormones [adrenal  
cortical stimulating hormone (ACTH) and  
thyroid stimulating hormone (TSH) are  
just two examples]
- other lymphokines and cytokines for  
systemic therapy
- interferon  $\gamma$  - for granulomatous disease  
of childhood (and other diseases being  
investigated)
- interferon  $\alpha$  - for leukemia and chronic  
active hepatitis
- erythropoietin - for chronic renal  
failure and other marrow suppressive  
disorders
- other hematologic growth factors - for  
marrow suppressive disorders
- administration e.g., tissue plasminogen  
activator for prevention of thrombosis  
in the pulmonary coronary arteries  
following reperfusion therapy,  
especially after balloon  
catheterization, or CD4 for human  
immunodeficiency virus (HIV) infection,  
and other recombinant proteins  
requiring systemic administration,  
whether short term or long term

- recombinant proteins for other hereditary disorders such as cerebrosidase deficiency and adenosine deaminase deficiency
- 5       • receptor agonists or antagonists - for example, for the control of systemic hypertension; interleukin-1 receptor antagonist for septic shock, rheumatoid arthritis and other disorders
- 10       • binding proteins for cytokines, lymphokines, and hormones - for example, tumor necrosis factor binding protein (a portion of the tumor
- 15       treatment of shock and wasting disorders mediated by tumor necrosis factor

For hereditary and acquired disorders of the gastrointestinal tract, this approach  
20 provides a means of administering recombinant proteins to the surface or within the cells or extracellular matrix of the walls of the gastrointestinal tract. Examples of possible applications include:

- 25       • pancreatic enzymes for pancreatic deficiency disorders such as cystic fibrosis
- lactase for lactose intolerance and the appropriate enzymes for the small
- 30       intestine disaccharidase deficiencies

- local therapy for gastrointestinal cancers with cytokines, tumor suppressor proteins (for example, p53 and retinoblastoma genes), and cytotoxic proteins
- prevention of cancer in individuals prone to gastrointestinal tract cancer (e.g., familial polyposis) with tumor suppressor proteins (for example, p53 and retinoblastoma genes).

For mammals and birds (more specifically, farm animals, for example - pigs, cattle, sheep, horses, dogs, cats, and chickens), this approach provides a means of administering recombinant proteins (for example, growth hormone) to these animals for the purposes of augmenting growth, generating characteristics for commercial purposes, and/or for general therapeutic purposes and for producing proteins from purified fractions given to humans, as well as antibodies for reagents reactive with human protein.

The use of proteins and polypeptides as therapeutic agents is greatly expanded according to the present invention by providing a means for delivering effective amounts of biologically active protein to a recipient individual. The preparations of this invention are suitably administered to animals, which include but are not limited to mammals (including humans), fish, and avians. The preparations are preferably administered to livestock (including cattle, horses, swine, sheep, goats, etc.), household pets (cats, dogs, canaries, parakeets, etc.) fish (especially in an aquarium or aquaculture environment, e.g., tropical fish, goldfish and

other ornamental carp, catfish, trout, salmon, etc.) and avians, especially poultry such as chickens, ducks, geese, etc.

In one embodiment of the present invention, the replication-deficient adenovirus can be employed with animal feeds (or, with less dosage control, with animal drinking water) acting as a nontoxic, pharmaceutically acceptable carrier for administration to animals, e.g., livestock, household pets, fish, poultry, etc. In one aspect, this embodiment is useful for producing proteins for purification to human sera and to produce antibodies for reagents against human proteins in a different species, e.g., cattle, horse, sheep, goat, rabbit, swine, etc. In another aspect, this embodiment is useful in the treatment of disorders in which proteins or polypeptides are useful therapeutic agents, particularly when the gene coding the therapeutic protein is derived from the species being treated, or with sequences which are closely homologous to prevent immune reactions.

The replication-deficient adenovirus of the present invention can likewise be employed in admixture with conventional excipients, i.e., pharmaceutically acceptable organic or inorganic carrier substances suitable for enteral (e.g., oral) application that do not deleteriously react with the virus. Suitable pharmaceutically acceptable carriers are well known in the art. (Suitable vehicles include those that are acid resistant and base sensitive, that is, sufficiently so such that transport can be effected through the stomach without unacceptable degradation.) They include but are not limited to water, salt solutions, alcohols, gum arabic, vegetable oils, benzyl alcohols, polyethylene

glycols, gelatine, carbohydrates such as lactose, anylose or starch, magnesium stearate, talc, silicic acid, viscous paraffin, perfume oil, fatty acid, monoglycerides and diglycerides, pentacrythritol fatty acid esters, hydroxy methyl cellulose, polyvinyl pyrrolidone, etc. Taking appropriate precautions not to kill the replication-deficient adenovirus, the preparations can be sterilized and if desired mixed with auxiliary agents, e.g., lubricants, preservatives, stabilizers, wetting agents, emulsifiers, salts for influencing osmotic pressure, buffers, coloring, flavoring and/or aromatic substances and the like that do not deleteriously react with the active virus. For example, agents can be used to increase the pH of the stomach, either directly (such as by buffers or bases) or indirectly (such as by drugs), to allow the virus to more readily pass through unharmed. They also can be combined where desired with other biologically active agents, e.g., antisense DNA or mRNA.

In another embodiment of the present invention, the replication-deficient adenovirus can be employed as a vaccine to develop immunity against infectious agents. The strategy is as follows. The gene coding for the protein against which immunity is to be developed is cloned into the replication deficient adenovirus construct; next, the replication deficient adenovirus containing the gene of interest is then administered to an animal (preferably, a human) as described herein. The gene sequences are designed such that the protein is secreted by the epithelial cells of the gastrointestinal tract into the systemic circulation such that immunity against the foreign proteins is developed.

Examples of this approach include the development of immunity against the hepatitis viruses, human immunodeficiency virus, and all other viruses that cause animal disease (particularly human disease). This strategy may also be used to develop immunity against bacteria, fungi, and other infectious agents.

A particularly interesting aspect of the present invention involves the use of replication-deficient adenovirus as a delivery system for chemotherapeutic agents, including antisense compounds, especially for use in cancer chemotherapy. Use with conventional chemotherapeutic agents is as discussed above. Briefly, use with antisense compounds for use against tumor cells in the bowel involves selecting mRNA as the primary drug target, with either another mRNA molecule or a synthetic oligodeoxynucleotide having the complementary base sequence to the mRNA forming a hybrid duplex by hydrogen-bonded base pairing. This hybridization can prevent expression of the target mRNA's protein product, a process called "translation arrest". Inhibition of mRNA is more efficient than inhibition of an enzyme active site because a single mRNA molecule gives rise to multiple protein copies. Thus, the selective inhibition of expression of a gene product required for cellular function yields the elusive but highly desired goal of chemotherapy: selective cell death. Such approaches are known in the literature, e.g., see J.S. Cohen, "Antisense Oligonucleotides as an Approach Toward Anti-Aids Therapy" at pages 195-224 in Design of Anti-Aids Drugs, E. deClerg (Ed), Elsevier Publishing Co. (1990); and S.L. Loke, et al. Current Topics in Microbiology and Immunology 141: 282-289 (1988).

For external applications, particularly suitable are tablets, dragees, liquids, drops, suppositories, or capsules. A syrup, elixir, or the like can be used wherein a sweetened vehicle is employed. Sustained or directed release compositions can be formulated, e.g., liposomes or those wherein the active virus is protected with differentially degradable coatings, e.g., by microencapsulation, multiple coatings, etc. It is also possible to freeze-dry the compositions and use the lyophilizates obtained.

Generally, the preparations of this invention are dispensed in unit dosage form comprising  $10^6 - 10^{14}$  pfu/ml of the replication-deficient adenovirus in a pharmaceutically acceptable carrier per unit dosage, preferably about  $10^{10} - 10^{12}$  pfu/ml. The dosages of the biologically active compounds administered according to this invention are generally known in the art but will frequently be reduced because of the improved delivery system provided by the present invention.

The actual preferred amounts of replication-deficient adenovirus administered in a specific case will vary according with the specific protein or polypeptide being utilized, the particular compositions formulated, the mode of application, and the particular situs and organism being treated.

The particular formulation employed will be selected according to conventional knowledge depending on the properties of the protein or polypeptide and the desired site of action to ensure bioavailability of the active ingredients, i.e., the extent to which the drug reaches its site of action or a biological fluid from which the drug has access to its site of action.



Dosages for a given host can be determined using conventional considerations, e.g., by customary comparison of the differential activities of the subject preparations and a known appropriate, conventional pharmacological protocol.

The present invention is described in further detail in the following non-limiting examples.

#### EXAMPLES

To demonstrate the feasibility of the invention, the replication deficient recombinant adenovirus Ad- $\alpha$ 1AT (Figure 1) was used to transfer the sequences coding for human  $\alpha$ 1-antitrypsin to the epithelial cells of the colon. Three models were used: (1) T84 human colon carcinoma cells in vitro; (2) intact rat colon ex vivo; and (3) cotton rat colon in vivo.

The following protocols and experimental details are referenced in the Examples that follow:

The recombinant vector (Ad- $\alpha$ 1AT) is constructed by deleting the majority of the E3 region and 2.6 mu from the left end of Ad5, and adding to the left end the  $\alpha$ 1-antitrypsin ( $\alpha$ 1AT) expression cassette from the plasmid pMLP- $\alpha$ 1AT, containing regulatory sequences and a recombinant human  $\alpha$ 1AT gene (Figure 1). Ad5 is commercially available from the American Type Culture Collection, Rockville, Maryland, USA. The methods for generating the  $\alpha$ 1AT cDNA, the expression cassette, and the final vector adenovirus are prepared using the methods described by M. Rosenfeld, et al. in Science 252: 431-434 (1991).

The bottom of Figure 1 presents details of the  $\alpha$ 1AT expression cassette. ITR, inverted ter-

5        minal repeat. To construct the recombinant viral  
vector Ad- $\alpha$ 1AT, the expression cassette was  
ligated with *Cla*I pre-cut Ad-d1327 DNA (to remove  
a portion of the E1a region from Ad-d1327). The  
10        recombinant adenovirus DNA was transfected into  
the 293 cell line where it was replicated,  
encapsulated into an infectious virus, and  
isolated by plaque purification. Individual  
plaques were amplified by propagation in 293  
15        cells and viral DNA extracted. The intactness of  
the DNA of the recombinant virus was confirmed  
prior to use by restriction fragment analysis and  
Southern hybridization. Stocks of Ad- $\alpha$ 1AT were  
propagated and titered in 293 cells. The virus  
20        was released from infected cells 36 hours post-  
infection by 5 cycles of freeze/thawing. The Ad-  
 $\alpha$ 1AT was further purified using CsCl gradients  
(for further details see Rosenfeld M et al.  
(1991) Science 252:431-434).

20        Figure 3A presents a demonstration of  
de novo synthesis and secretion of human  $\alpha$ 1-  
antitrypsin ( $\alpha$ 1AT) by rat colon exposed to the  
recombinant adenovirus Ad- $\alpha$ 1AT. The colon was  
washed, the end tied off to make a "sausage"  
25        2 - 5 cm. in length, and 50 - 100 microliters of  
 $10^{10}$  -  $10^{12}$  pfu/ml Ad- $\alpha$ 1AT in LHC-8 medium  
injected into the lumen. The "sausage" was then  
incubated for 24 hr. 37°, washed, cut into 1mm<sup>2</sup>  
fragments, and <sup>32</sup>S-methionine (500 mCi/ml) added  
30        in methionine-minus LHC-8 medium. After incu-  
bation for 24 hr, 37°, the fluid bathing the  
fragments were evaluated for the presence of  
human  $\alpha$ 1AT by immunoprecipitation, sodium  
dodecyl sulfate acrylamide gels and  
35        autoradiography. The results are set forth in  
Figure 3B, wherein: Lane 1 - uninfected  
colon; lane 2 - colon infected with Ad- $\alpha$ 1AT; and  
lane 3 - same as lane 2, but with the antibody

exposed to unlabeled human  $\alpha$ 1AT (to demonstrate the specificity of the antibody). The 52 kDa human  $\alpha$ 1AT is indicated by the arrow.

#### EXAMPLE 1

##### 5      T84 Human Colon Carcinoma Cells In Vitro

2      This model was used to demonstrate that  
3      human colon epithelial cells can be infected by  
4      Ad- $\alpha$ 1AT, and that the infection resulted in the  
5      secretion of human  $\alpha$ 1AT to the apical surface  
6      (i.e., to the lumen side of the epithelium) and  
7      to the basolateral surface (i.e., the circulation  
8      side of the epithelium). To accomplish this, the  
9      T84 cell line was grown on microporous membranes  
10     until they became confluent and formed tight  
11     junctions (electrical resistance  $>150 \text{ ohm-cm}^2$   
12     across the epithelium). The microporous  
13     polycarbonate membrane ( $4.7 \text{ cm}^2$ , pore size  $3.0 \text{ um}$ ,  
14     Transwell Col., Coster, Cambridge, MA) with  
15     the epithelial cells separate two chambers that  
16     contain culture fluid, i.e., an in vitro system  
17     that mimics the epithelium in vivo. The upper  
18     chamber faces the apical surface and the lower  
19     chamber faces the basolateral surface. The  
20     combination of the cells and the tight junctions  
21     between the cells physically separate the fluids  
22     and the upper and lower chambers (equivalent to  
23     the in vivo situation where the apical surface  
24     abuts the inside lumen of the colon and the  
25     basolateral surface abuts the tissue side (and  
26     thus the circulation; see Figure 2). The cells  
27     are cultured in DMEM, 2% fetal calf serum for 1.5  
28     hr,  $37^\circ$  and then in DMEM, 10% fetal calf serum  
29     for 24 hr,  $37^\circ$  with no adenovirus or with Ad- $\alpha$ 1AT  
30     (from the apical side as would occur in vivo).  
31     Three different intensities of infection were  
32     used [measured in plaque-forming units (pfu), the

number of infectious viral particles per ml of fluid;  $5 \times 10^9$ ,  $10^{10}$ , and  $2.5 \times 10^{10}$  pfu/culture]. The media was then collected and evaluated for the presence of human  $\alpha 1$ -antitrypsin using an enzyme-linked immunoassay (Wewers MD et al. (1987) N Engl J Med 316: 1055-1062). The data demonstrates that Ad- $\alpha 1$ AT infection causes the human colon epithelial cells to secrete  $\alpha 1$ AT, and to do so in both directions, i.e., to the apical and basolateral surfaces. The amount secreted to the apical surface compared to the basolateral surface ranged from 3.98 to 4.69 (average 4.34) i.e., for every 4.34 molecules secreted into the lumen (where it would eventually be excreted in vivo), 1 molecule would be secreted into the tissue (where it would be available to the circulation).

The results of Example 1 are shown in Table I.

20

Table I

Polarity of Secretion of Human  $\alpha 1$ -Antitrypsin by the Human T84 Colon Carcinoma Cell Line Following Infection by the Recombinant Adenovirus Ad- $\alpha 1$ AT

Infection <sup>1</sup>	Amount $\alpha 1$ AT Secreted in 24 hr ( $\mu$ g) <sup>2</sup>		Ratio of $\alpha 1$ AT in Apical and Basolateral Compartments
	Apical	Basolateral	
None	0	0	---
$5 \times 10^9$	3.54	0.89	3.98
$10^{10}$	7.42	1.58	4.69
$2.5 \times 10^{10}$	8.89	2.05	4.34

<sup>1</sup> number of pfu Ad- $\alpha 1$ AT added to the culture; all cells were grown on 4.7 cm<sup>2</sup> microporous membranes until tight junctions were formed (electrical resistance > 150 ohm-cm<sup>2</sup>)

<sup>2</sup> Measured by enzyme linked immunoassay (Wewers MD et al. (1987) N Engl J Med 316: 1055-1062)

EXAMPLE 2Intact Rat and Cotton Rat Colon In Vitro

This model was used to determine if the recombinant adenovirus can infect colon epithelial cells in circumstance where the cells were normal (i.e., not derived from a neoplasm as in the T84 model) and were in their normal architectural configuration. To do this, rat colon was removed, washed and a 2-3 cm section made into a closed "sausage" by tying off both ends (Figure 3). Ad- $\alpha$ 1AT was injected into the lumen (e.g., equivalent to live recombinant adenovirus being released from enteric coated capsules). The "sausage" was placed in culture media for 24 hr, 37° and then evaluated in two ways.

First, the colon was fragmented into 1 mm<sup>3</sup> pieces, <sup>35</sup>S-methionine was added, the culture continued for 24 hr, 37°, and the ability of the colon to de novo synthesize and secrete human  $\alpha$ 1AT evaluated using immunoprecipitation, sodium dodecyl sulfate acrylamide gels and autoradiography (see Rosenfeld M et al. (1991) Science 252:431-434 for details of the methods). The results demonstrate that uninfected rat colon does not synthesize and secrete human  $\alpha$ 1AT in vitro, but that Ad- $\alpha$ 1AT infected rat colon does (Figure 3).

A similar technique was used to evaluate cotton rat colon, but using enzyme-linked immunoassay (ELISA) to quantify the amount of human  $\alpha$ 1AT secreted into the lumen (i.e., apical secretion; it is not possible to evaluate basolateral secretion in this model). Following 48 hr infection with approximately 10<sup>11</sup> pfu Ad- $\alpha$ 1AT injected into the lumen of cotton rat colon

"sausage" in vitro, evaluation of the luminal fluid demonstrated  $3.3 \pm 0.6 \mu\text{g/ml}$  human  $\alpha\text{1AT}$ .

### EXAMPLE 3

#### Cotton Rat Colon In Vivo

5           This model was used to demonstrate that the concept will work in vivo in living animals. Two strategies were used, both in cotton rats. First, following general anesthesia and laparotomy, a section of colon was ligated in two  
10 places to form an in vivo "sausage" in a fashion that permitted normal blood flow to that segment. Ad- $\alpha\text{1AT}$  was injected into the lumen and the laparotomy closed. The animals were maintained without oral intake. After 48 hr, a serum sample  
15 was taken and evaluated for the presence of human  $\alpha\text{1AT}$  by ELISA. Second, following general anesthesia and laparotomy  $10^{10} - 10^{12}$  pfu of, Ad- $\alpha\text{1AT}$  was injected into the lumen of the colon without ligation. After 48 hr, a serum sample  
20 was taken and evaluated for human  $\alpha\text{1AT}$  by ELISA. In both cases human  $\alpha\text{1AT}$  was clearly evident.

The results are shown in Table II.

Table II

25   Serum Levels of Human  $\alpha\text{1-antitrypsin}$  in Cotton Rats 48 hr Following In Vivo Administration of Ad- $\alpha\text{1AT}$  in the Lumen of the Colon

Condition	Serum $\alpha\text{1AT}$ Level (ng/ml) <sup>3</sup>
Mock infection	0
"Sausage" infection <sup>1</sup>	$145 \pm 29$
Direct infection <sup>2</sup>	$74 \pm 9$

<sup>1</sup> Ad- $\alpha\text{1AT}$  (50-100  $\mu\text{l}$ , approximately  $10^{11}$  pfu) injected into the lumen of a segment of bowel isolated by ligation at both ends.

30   <sup>2</sup> Similar to the "sausage" infection, but without isolating a segment of bowel by ligation.

<sup>3</sup> Measured by enzyme linked immunoassay (Wewers MD et al. (1987) N Engl J Med 316: 1055-1062).

While the foregoing invention has been described in some detail for purposes of clarity and understanding, it will be appreciated by one skilled in the art from a reading of this disclosure that various changes in form and detail can be made without departing from the true scope of the invention and appended claims.

WHAT IS CLAIMED IS:

1. A method of producing a biologically active protein in the gastrointestinal tract of an individual, comprising:

administering to said individual's gastrointestinal tract a replication deficient adenovirus comprising a DNA segment encoding said protein under conditions such that said protein is produced.

2. The method according to claim 1, wherein said protein is a therapeutic protein.

3. The method according to claim 1, wherein said protein is selected from the group consisting of a coagulation factor, a pituitary hormone, a peptide hormone, a lymphokine, a cytokine, a tumor suppressor protein, a hematologic growth factor, a receptor agonist, and a receptor antagonist.

4. The method according to claim 1, wherein said protein is selected from the group consisting of  $\alpha$ 1-antitrypsin, erythropoietin, Factor VIII, growth hormone, tumor necrosis binding protein, interleukin-1 receptor antagonist, interferon  $\gamma$ , interferon  $\alpha$ , and insulin.

5. The method according to claim 1, wherein said adenovirus is Ad- $\alpha$ 1AT.

6. The method according to claim 1, wherein said adenovirus is administered in an enteric capsule.



7. An enteric capsule comprising a replication deficient adenovirus containing a DNA segment encoding a therapeutic protein.

8. A method of producing a biologically active protein in the gastrointestinal tract of an animal, comprising:

administering to the gastrointestinal tract of an animal a replication deficient adenovirus comprising a DNA segment encoding said protein in an amount effective and under conditions such that said protein is produced.

9. The method according to claim 8, wherein said animal is a mammal, avian or fish.

10. The method according to claim 9, wherein said animal is selected from the group consisting of pig, sheep, cattle, horse, cat, and dog.

11. The method according to claim 9, wherein said animal is a chicken.

12. A pharmaceutical composition comprising:  
a replication deficient adenovirus containing at least one DNA segment encoding for a therapeutic protein, said adenovirus contained in a vehicle that is acid-resistant and base-sensitive and  
a pharmaceutically acceptable diluent, carrier, or excipient.

13. A method of developing immunity against a protein in an animal comprising:

administering to said animal's gastrointestinal tract a replication

deficient adenovirus comprising a DNA segment encoding said protein in an amount effective and under conditions such that said protein is produced and said immunity against said protein is developed.

## ADENOVIRUS

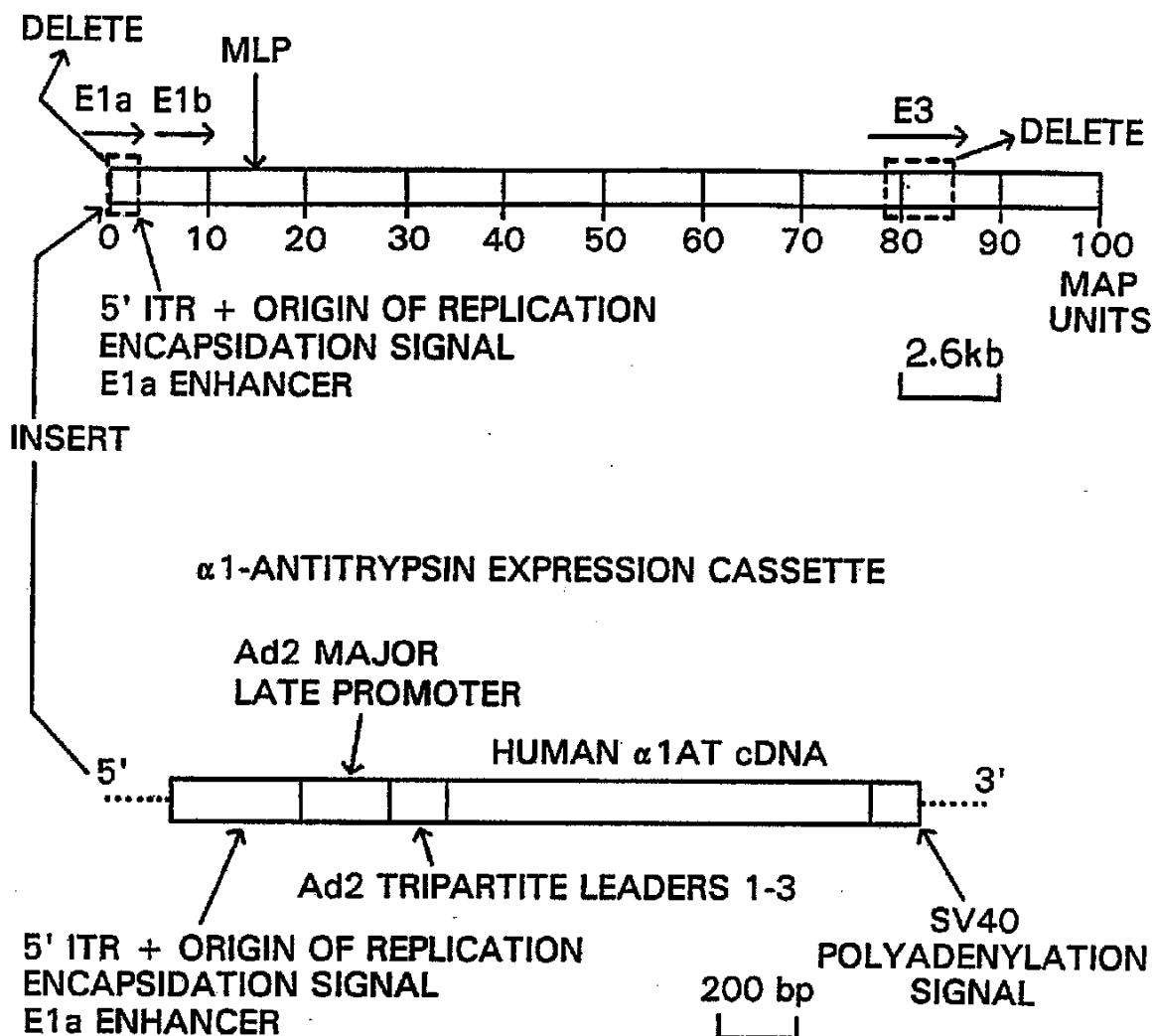


FIG. 1

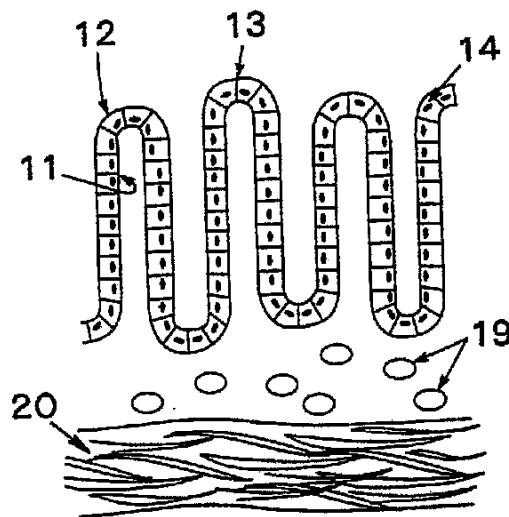


FIG. 2A

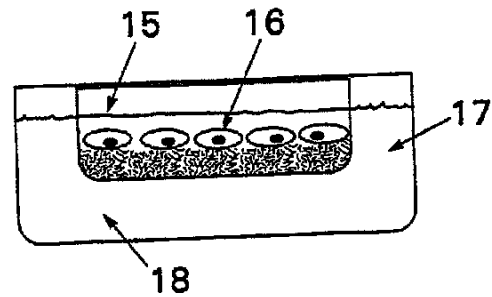


FIG. 2B

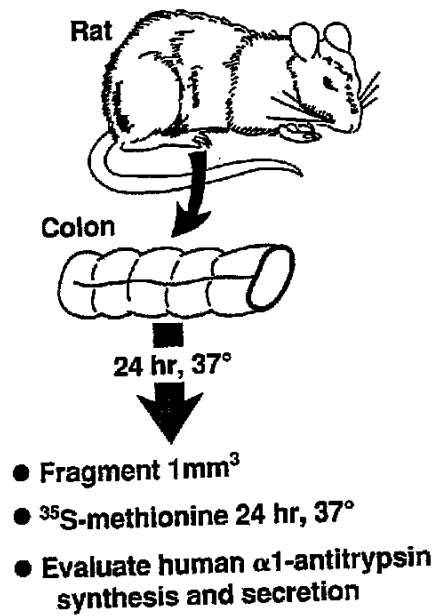


FIG. 3A

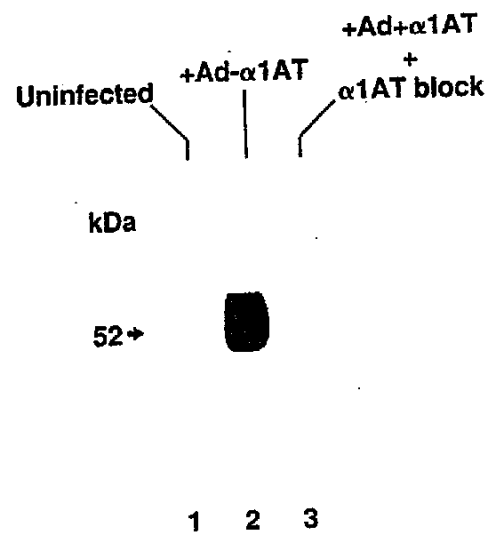


FIG. 3B

## INTERNATIONAL SEARCH REPORT

International application No.  
PCT/US92/07029

**A. CLASSIFICATION OF SUBJECT MATTER**

IPC(5) : A61K 48/00; C12N 15/87, 15/63, 15/57, 15/12

US CL : 424/93A; 435/69.1, 69.2, 69.3, 69.4, 69.5, 69.6, 320.1; 935/32, 62

According to International Patent Classification (IPC) or to both national classification and IPC

**B. FIELDS SEARCHED**

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 424/93A; 435/69.1, 69.2, 69.3, 69.4, 69.5, 69.6, 320.1; 935/32, 62

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

BIOSIS, MEDLINE, APS

**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	US, A, 4,920,209 (DAVIS ET AL.) 24 April 1990, see the entire document.	1-13
Y	US, A, 4,980,286 (MORGAN ET AL.) 25 December 1990, see the entire document.	1-13
Y	Science, Volume 252, issued 19 April 1991, Rosenfeld et al., "Adenovirus-Mediated Transfer of a Recombinant $\alpha$ 1-Antitrypsin Gene to the Lung Epithelium in Vivo", pages 431-434, see the entire document.	1-13
A	Biotechniques 6, issued 1989, Berkner, "Development of Adenovirus Vectors for the Expression of Heterologous Genes", pages 616-629, see the entire document.	1-13
A	FEBS Letters, Volume 267, number 1, issued July 1990, Gilardi et al., "Expression of human $\alpha$ 1-antitrypsin using a recombinant adenovirus vector", pages 60-62, see the entire document.	1-13

☐ Further documents are listed in the continuation of Box C. ☐ See patent family annex.

* Special categories of cited documents:	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"A" document defining the general state of the art which is not considered to be part of particular relevance	"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"E" earlier document published on or after the international filing date	"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"L" document which may throw doubt on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"Z" document member of the same patent family
"O" document referring to an oral disclosure, use, exhibition or other means	
"P" document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search 05 November 1992	Date of mailing of the international search report 3 NOV 1992
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